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1 **Fibromyalgia: Heterogeneity in personality and psychopathology and its**
2 **implications**

3 **Abstract**

4 The fibromyalgia syndrome (FM) is a chronic widespread pain condition whose
5 etiology remains unknown and no treatment has satisfactory levels of success. A meta-
6 analysis has identified a distinct Minnesota Multiphasic Personality Inventory-2
7 (MMPI-2) clinical profile between FM female patients and healthy controls, and
8 differences between FM and other chronic pain condition with clear etiology have also
9 been found. However, heterogeneity in this population has been suggested in several
10 studies. We aim to assess clinical aspects in FM patients, based on personality
11 psychopathology characteristics, in order to explore heterogeneity and the existence of
12 core common aspects. In this cross-sectional study, a relatively homogeneous sample of
13 56 female FM patients ($M_{age} = 45.95$, $SD_{age} = 9.39$) was assessed through MMPI-2. A
14 K-Means cluster analysis identified two clusters, one ($n = 24$) with clinically significant
15 levels in *Negative Emotionality* and *Introversion* scales. Subsequent MANOVAs
16 identified important features of this cluster on several MMPI-2 dimensions. Moreover,
17 several dimensions are clinically elevated in both clusters. In conclusion, the
18 combination of psychopathological negative emotionality, interpersonal isolation, and
19 low hedonic capacity, in a group of patients, has implications for the daily living and
20 treatment of FM patients, and several core aspects of FM need to be addressed.

21 **Keywords:** Fibromyalgia; personality characteristics; psychopathology dimensions;
22 MMPI-2

23 Fibromyalgia (FM) is a syndrome characterized by chronic widespread pain,
24 which is frequently associated with fatigue, sleep disorder, other functional somatic
25 syndromes, mental and physical disorders, as well as disability and diminished quality

1 of life. Although a central sensitization phenomena has been associated to FM (Ablin et
2 al., 2012; Arnold et al., 2016) the etiopathology of FM remains unknown (Thieme,
3 Mathys, & Turk, 2017). Some authors conceptualize this syndrome as part of a group of
4 affective spectrum disorders (e.g., Arnold et al., 2004), as it has a high comorbidity with
5 psychiatric disorders. In addition, a high prevalence of alexithymia has been found in
6 FM patients (Di Tella et al., 2018). The fact that FM is a medically unexplained
7 syndrome, and the quality of evidence of a large range of treatments for FM are only
8 modest and has not shown significant improvement over the past two decades (Thieme
9 et al., 2017) presents a challenge to clinical psychology and its contribution to the
10 understanding and treatment of FM.

11 In the scope of an integrative biopsychosocial approach to FM, psychological
12 aspects may play an important role as predisposing factors to FM, and personality is one
13 of those aspects (Eich, Hartmann, Muller, & Fischer, 2000; Malin & Littlejohn, 2012;
14 Van Houdenhove, Luyten, & Egle, 2009). Within a diathesis-stress model of disease,
15 some personality and psychopathology features that make people more vulnerable to
16 stressors would be antecedent to FM, interacting with physiological vulnerabilities to
17 the development of the syndrome (Thiagarajah Guymer, Leech, & Littlejohn, 2014).

18 A recent meta-analysis focused on the Minnesota Multiphasic Personality
19 Inventory (MMPI-2) has shown that female FM patients have a psychopathology profile
20 significantly different than the profile of healthy volunteers (Novo, Gonzalez, Peres, &
21 Aguiar, 2017a; 2017b). Nevertheless, it also acknowledged that the FM patients are
22 probably a heterogeneous group regarding personality and psychopathology profiles.

23 Pertaining to psychopathology, with the original MMPI, Ahles, Yunus, Riley,
24 Bradley, & Masi (1984) found a large group of FM patients with a clinical profile
25 within the normal range, followed by a group with the typical chronic pain profile

1 (clinically significant scores on the “neurotic triad scales”, i.e., hypochondriasis,
2 depression and hysteria), and a smaller group with the psychopathological profile
3 (significant elevations in several clinical scales) was found. On the contrary, some
4 studies found a larger cluster with the psychopathological profile (Claros et al., 2006;
5 Porter-Moffitt et al., 2006).

6 Pertaining to personality only, within the five factor model of personality, a
7 cluster with higher Neuroticism and lower Conscientiousness is identified (Bucourt et
8 al., 2018; Torres et al., 2013) and also with lower Extraversion (Torres et al., 2013)
9 associated with more self-reported pain (Bucourt et al., 2018).

10 Finally, the distressed type or Type D personality, a specific combination of
11 Negative Affectivity and Social Inhibition, has been more recently studied in this
12 population, and it constitutes our main interest. Van Middendorp et al. (2016) found a
13 high prevalence within an FM sample, and Ablin, Zohar, Zaraya-Blum, and Buskila
14 (2016) found a higher prevalence of Type D personality in one of the FM clusters,
15 associated with a less adaptive pattern.

16 In conclusion, we intend to explore heterogeneity based on structural personality
17 psychopathology dimensions (as most of the studies focus on clinical psychopathology
18 dimensions), within a community sample with a homogeneous age range. We
19 hypothesize that different clinical levels in the psychopathology and personality
20 dimensions will be found.

21 **Method**

22 ***Participants***

23 The participants were 56 FM female patients between 30 and 60 years old ($M =$
24 45.95 ; $SD = 9.39$). The inclusion criteria were: Having a pure FM diagnosis (not having
25 another rheumatic disease or painful condition) for at least six months. Finally, all the

1 participants had the response consistency levels (VRIN and TRIN validity scales)
2 within the normal range.

3 ***Instruments***

4 We used MMPI-2 (Butcher, Dahlstrom, Graham, Tellegen, & Kaemmer, 1989) in
5 a Portuguese version (Silva, Novo, Prazeres, & Pires, 2006), is a self-administered
6 inventory to assess clinical and personality psychopathology. The results are converted
7 into normalized T-scores, and in general terms, $T > 65$ are clinically significant. In
8 clinical scales, internal consistency coefficients range between .34 and .87, most of them
9 in .80.

10 ***Procedure***

11 Most of the FM patients were recruited through a patient`s association, contacted by
12 telephone and asked about the inclusion criteria and availability to participate in the
13 study. The remaining FM patients were recruited within the scope of a psychological
14 assessment service in a university centre open to the community. Research with MMPI-
15 2 was approved by the ethic committee of the North Lisbon Hospital Center. The
16 informed consent was obtained and the privacy was observed in accordance with the
17 principles of the Declaration of Helsinki.

18 ***Data analysis***

19 We conducted K-means non-hierarchical cluster analysis, based on the five
20 MMPI-2 personality psychopathology scales (PSY-5: *Agressiveness, Psychoticism,*
21 *Disconstraint, Negative Emotionality/Neuroticism, and Introversion/Low Positive*
22 *Emotionality*) testing a three clusters solution in the first place. As it was not
23 appropriate, we tested a two clusters solution. We used Chi-Square test to identify the
24 differences between the clusters in the sociodemographic nominal and ordinal variables
25 and a one-way analysis of variance (ANOVA) to test the differences in age and

1 diagnosis duration. We used multivariate analysis of variance (MANOVA) to test the
2 differences between the two clusters in the MMPI-2 clinical scales.

3 **Results**

4 In the K-means two-cluster solution, based on the scores of the PSY-5 MMPI-2
5 scales, convergence was achieved after three iterations. The distance between final
6 cluster centers was 23.945. The characterization features are presented in Table 1.

7 The larger cluster ($n = 32$) has no clinically significant elevation, and the other
8 cluster ($n = 24$) has clinically significant elevations ($T \geq 65$) in two scales: *Negative*
9 *Emotionality* ($M = 65.25$, $SD = 9.66$; $t = -3.38$; $p = .001$) and *Introversion* ($M = 68.38$,
10 $SD = 9.90$; $t = -8.61$; $p = .000$) (Figure 1).

11 The MANOVA showed significant differences between the clusters in the
12 composite of clinical scales (Wilk's $\Lambda = .415$; $F(10,45) = 6.344$; $p = .000$; $\eta^2_{par} = .585$).
13 Both clusters had clinically significant elevations in *Hypochondriasis*, *Depression*,
14 *Hysteria*, and *Schizophrenia* (Figure 2), and Cluster 2 also had it in *Psychasthenia* ($M =$
15 69.63 , $SD = 8.99$; $t = -3.54$; $p = .001$).

16 **Discussion**

17 As hypothesized, we found different clinical levels in the psychopathology and
18 personality dimensions of FM patients, organized in two clusters. One of them (cluster
19 2) is in line with Type D personality found by Ablin et al. (2016), and it is relevant that
20 this cluster has higher *Introversion* mean levels than the *Neuroticism* ones. As the
21 presence of positive affect protects against the experiencing of negative affect in times
22 of stress and pain (Davis, Zautra, & Smith, 2004), this protective aspect is absent in this
23 cluster. As Vendrig, Derksen, & Mey (2000) found that pretreatment higher scores in
24 *Introversion* predicted decreased satisfaction with the treatment, decreased self-
25 perceived emotional change, and a probability of investing less energy in the treatment
26 process, this could be a relevant implication to any intervention with these patients.

1 Cluster 2 has an extremely high mean level of *Depression*, as the most elevated
2 scale of the neurotic triad, a configuration that does not characterize chronic pain
3 patients in general, as *Hypochondriasis* and *Hysteria* are usually the more elevated
4 scales in these patients (Ahles et al., 1984; Claros et al., 2006; Porter-Moffitt et al.,
5 2006; Keller & Butcher, 1991). Cluster 2 has also clinically significant levels of
6 *Psychasthenia*, which corresponds to a diagnosis close to Obsessive-Compulsive
7 Disorder (Graham, 2012).

8 Apart from heterogeneity, we identified common features of the FM patients.
9 Both clusters have clinically significant levels in the neurotic triad, and in
10 *Schizophrenia*, reflecting a pathological clinical profile, predominantly neurotic but also
11 reflecting social alienation, unusual beliefs, confusion and lack of adequacy, which
12 differentiates FM from other chronic pain samples, in which the neurotic triad only is
13 expected to be elevated (e.g., Keller & Butcher, 1991).

14 A group of FM patients have relevant personality, clinical and specific features
15 that undoubtedly may compromise any regular chronic pain treatment. This personality
16 pattern has a double implication, at both emotional and relational level, as high
17 Negative Emotionality hinders relations with others, and high Introversion leads to
18 emotional and social disengagement (Friedman, Lewak, Nichols, & Webb, 2001).
19 Beyond heterogeneity, affective distress is a common aspect to the sample and it should
20 require psychological intervention as part of any medical intervention.

21 This work has some limitations, mainly the sample dimension, and the absence of
22 a control group. The fact that the sample is composed by women only makes the
23 interpretations appropriate to female fibromyalgia patients only. As strengths of the
24 study, we aimed for a FM community sample, with a homogeneous age range and, to
25 our best knowledge, it is also the first study exploring the heterogeneity in the clinical

1 and psychopathological symptomatology in FM patients, based exclusively on structural
2 and relatively stable personality characteristics.

3 In future research it would be important to study the adherence of FM patients to
4 medical and psychological interventions, the treatment results, and relate them to the
5 personality and psychopathology features.

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